



A drug discovery company dedicated to addressing unmet medical needs

Focusing on the Central nervous system (CNS) and
Autoimmune systems

September 2024
Company Presentation

Disclaimer

The information contained in this presentation and any accompanying oral presentation ("Presentation") is being supplied to you by Argent BioPharma on behalf of itself and its subsidiaries (together, "Argent BioPharma"). By accepting this Presentation, you agree to be bound by the following terms.

The content of this Presentation has not been approved by an authorised person within the meaning of the Financial Services and Markets Act 2000 (FSMA). This Presentation is for background purposes only and is not intended to be relied upon as advice to investors or potential investors, and does not contain all information relevant or necessary for an investment decision. The Presentation should be read in conjunction with Argent BioPharma's other periodic and continuous disclosure announcements filed with the Australian Securities Exchange. This Presentation does not constitute or form part of, and should not be construed as, an offer for sale or subscription of, or solicitation of any offer to buy or subscribe for, any securities of Argent BioPharma nor should it or any part of it form the basis of, or be relied on in connection with, or act as an inducement to enter into any contract or commitment whatsoever. The merits and suitability of any investment action in relation to securities should be considered carefully and involve, among other things, an assessment of the legal, tax, accounting, regulatory, financial, credit and other related aspects of such securities. This Presentation is exempt from the general restriction set out in section 21 of FSMA on the communication of financial promotions.

This communication is exempt under the Financial Services and Markets Act 2000 (Financial Promotion) Order 2005 (as amended) (the "Order") on the grounds that it is made or directed only to: (a) persons having professional experience in matters relating to investments within the definition of investment professionals in Article 19(5) of the Order; and (b) recipients who are high net worth bodies corporate, unincorporated associations, and partnerships and trustees of high value trusts as described in Article 49(2) of the Order; (all such persons together being referred to as "Relevant Persons"). Persons of any other description, including those that do not have professional experience in matters relating to investments, should not rely or act upon this Presentation, and the controlled investment and controlled activity to which this Presentation relates will only be available to Relevant Persons. If a person is unsure whether they would be classed as a Relevant Person, then they should seek independent legal advice or advice from an investment professional.

Not financial product advice

This document does not constitute financial product advice or take into account your investment objectives, taxation situation, financial situation or needs. This document consists of summarised information and does not involve or imply a recommendation of a statement of opinion in respect of whether to buy, sell or hold a financial product. An investment in Argent BioPharma is considered to be speculative in nature. Before making any investment decision in connection with any acquisition of securities, investors should consult their own legal, tax and/or financial advisers in relation to the information in, and action taken on the basis of, this document.

No Liability

The information contained in this Presentation has been prepared in good faith by Argent, however no guarantee, representation or warranty, expressed or implied is, or will be made, by any person (including Argent and its affiliates and their directors, officers, employees, associates, advisers and agents) as to the accuracy, reliability, correctness, completeness or adequacy of any statements, estimates, options, conclusions or other information contained in this document. To the maximum extent permitted by law, Argent BioPharma and its affiliates and their directors, officers, employees, associates, advisers and agents each expressly disclaims any and all liability, including, without limitation, any liability arising out of fault or negligence, for any loss arising from the

use of or reliance on information contained in this document including representations or warranties or in relation to the accuracy or completeness of the information, statements, opinions, forecasts, reports or other matters, express or implied, contained in, arising out of or derived from, or for omissions from, this document including, without limitation, any financial information, any estimates or projections and any other financial information derived therefrom. Statements in this document are made only as of the date of this document unless otherwise stated and the information in this document remains subject to change without notice.

Argent BioPharma, nor its related bodies corporate, officers, their advisers, agents and employees accept any responsibility or liability to you or to any other person or entity arising out of this Presentation including pursuant to the general law (whether for negligence, under statute or otherwise), pursuant to the general law (whether for negligence, under statute or otherwise), or under the Australian Securities and Investments Commission Act 2001, Corporations Act 2001, Competition and Consumer Act 2010 or any corresponding provision of any Australian state or territory legislation (or the law of any similar legislation in any other jurisdiction), or similar provision under any applicable law. Any such responsibility or liability is, to the maximum extent permitted by law, expressly disclaimed and excluded.

Forward looking statements

The information in this presentation is for general information only. To the extent that certain statements contained in this presentation may constitute "forward-looking statements" or statements about "future matters", the information reflects Argent BioPharma's intent, belief or expectations at the date of this presentation. In some cases forward looking statements can be identified by the use of terms such as "believes", "estimates", "anticipates", "projects", "expects", "intends", "may", "will", "seeks" or "should" or variations thereof, or by discussions of strategy, plans, objectives, goals, future events or intentions. Subject to any continuing obligations under applicable law or any relevant listing rules of the Australian Securities Exchange, Argent BioPharma disclaims any obligation or undertaking to provide you with access to any additional information or to update this Presentation or to correct any inaccuracies in, or omissions from this presentation which may become apparent.

Any forward-looking statements, including projections, guidance on future revenues, earnings and estimates, are provided as a general guide only and should not be relied upon as an indication or guarantee of future performance. Forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause Argent BioPharma's actual results, performance or achievements to differ materially from any future results, performance or achievements expressed or implied by these forward-looking statements. Any opinions, projections, estimates or forecasts contained in the Presentation constitute a judgment of Argent BioPharma only and should not be relied upon, and are provided as at the date of this Presentation and are subject to change without notice. This information is subject to change without notice. The accuracy of such information is not guaranteed, it may be incomplete or condensed and it may not contain all material information concerning Argent BioPharma and its subsidiaries. The forward-looking information contained in the Presentation is expressly qualified by this cautionary statement.

This Presentation must not be distributed, published, reproduced or otherwise made available to any person, in whole or in part, for any purposes whatsoever with addresses the United States, its territories or possessions or in any other jurisdiction outside of Australia or the United Kingdom where such distribution or availability may lead to a breach of any law or regulatory requirements. The distribution of this Presentation in other jurisdictions may be restricted by law, and persons into whose possession this Presentation come should inform themselves about, and observe, any such restrictions. Any failure to comply with these restrictions may constitute a violation of the laws of the relevant jurisdiction and you undertake to indemnify Argent for any loss or damage incurred as a result of your breach of this undertaking.

Argent BioPharma is a specialized Biopharmaceutical company focused on developing and marketing innovative drugs.

Committed to providing accessible medical treatments for unmet medical needs.

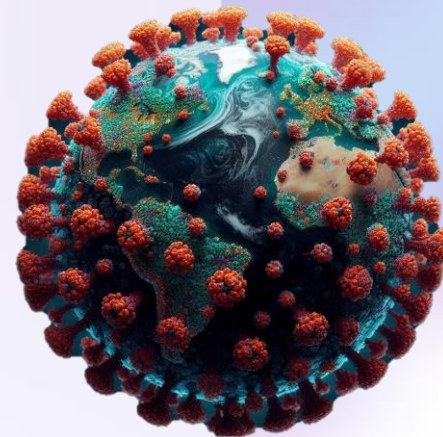
Innovative Technology Platform:

Utilizing advanced nanotechnology and a multidisciplinary approach in its in-house R&D platform to develop innovative therapies. This approach mainly targets the central nervous system (CNS) and the Autoimmune System

Argent BioPharma Vision and Mission

Vision: Argent BioPharma aspires to become a leading biopharmaceutical company recognized globally for its innovative treatments that address unmet medical needs, particularly in the central nervous system (CNS) and immune-related diseases. The company aims to be a key player in transforming patient outcomes by providing advanced, effective therapies for conditions with limited or no treatment options.

Mission: Argent BioPharma aims to improve patient outcomes by providing effective, cutting-edge treatments for conditions with limited or no existing therapeutic options. This involves leveraging its EU GMP manufacturing facilities, robust drug pipeline, and experienced management team to bring these innovative therapies to market as quickly and efficiently as possible.



The background features a light blue and white abstract design. On the left, there is a circular inset showing a close-up of a petri dish with several wells containing a dark, granular substance. This circle overlaps with a larger, semi-transparent purple circle. To the right of the purple circle is a large, bold text block. Thin, curved lines in light blue and white sweep across the background, adding a sense of motion and design.

R&D Centers and Manufacturing Capabilities

(3 sites in total)

In-house Research, Development & Production

Wholly owned and Operated EU GMP Sites

- **Slovenia, Emonska R&D hub** is staffed with over 40 highly experienced pharmaceutical industry professionals dedicated to developing advanced drug formulations and delivery systems.
- **Slovenia, Emonaka Quality and medical site**, expertly navigates regulatory pathways to secure marketing approvals from both the US FDA and EMA markets.
- **Slovenia, Kmaniska IMP EU GMP site**, where the formula is taken to production for clinical trials and early patient access.
- **Malta, Hal Far Fully automated EU GMP site**, possesses advanced production capabilities, enabling efficient manufacture of drugs and generating sales.



Slovenia - EU Research and Development site

Integrated research hub ensuring innovation, patient access, long-term value, and a strong market position.

- **Advanced R&D Operations:** Operates with a team of skilled professionals, ensuring continuous innovation and progress in drug development.
- **Early Patient Access:** Capable of developing and producing products for Early Patient Access, ensuring critical therapies reach patients swiftly.
- **Long-term Value:** The R&D center drives innovation, expanding Argent's drug portfolio and generating long-term value.
- **Investment and Market Position:** Ongoing innovation attracts investment, enhances market position, and addresses unmet medical needs, providing substantial returns to shareholders.



Malta Manufacturing site

Fully built EU GMP manufacturing facility

A modern EU GMP-certified manufacturing facility, operational for less than three years and supported by multi-million dollar grants from the Maltese government¹.

- **Automated Production Capabilities:** The facility features fully automated production capabilities, producing tens of thousands² of drug units per shift, ensuring high-volume, high-quality output.
- **Versatile Drug Manufacturing:** The facility supports Argent BioPharma's therapeutic portfolio by being equipped to manufacture a wide range of drugs, including flagship products like CannEpi[®] and CimetrA[®].

1. Refer to ASX Announcement dated 2 November 2021

2. Malta GMP certificate. Based on 30ml bottle size





Key Products

We utilize advanced nanotechnology and a multidisciplinary approach in our in-house R&D platform to develop innovative therapies for unmet medical needs.

Early-Stage Revenue Generating Refractory Epilepsy Seizure Control Treatment

- Approximately 30% of generalized seizure epilepsy patients have Refractory Epilepsy, aka “Drug-Resistant Epilepsy” (DRE)¹, representing 15,000,000 patients around the world.
- CannEpil® is now available to patients in the UK by **Named Patient Request**, to be prescribed by clinicians listed on the **GMC Specialist Register**.
- CannEpil® is accepted by the Irish Health Product Regulatory Authority (HPRA), and full health insurance coverage is obtained through the Medicinal Cannabis access program (MCAP)
- The first UK patient has access to CannEpil® through the **NHS RESCAS** pathway and the “I am Billy Foundation”, they have been using CannEpil® for over 1.5 years.



Total Addressable
Global Market: £15B²

1. Fattorusso A, et al. (2021) The Pharmacoresistant Epilepsy: An Overview on Existent and New Emerging Therapies. Front. Neurol.
2. <https://www.biospace.com/epilepsy-therapeutics-market-worth-15-1-bn-by-2030-at-a-cagr-of-4-5-percent>
Date on file – CannEpil Driving Performance
Data on file – CannEpil vs. MXP100 study

Market potential

Worldwide
Potential
Patients with DRE

15M

UK Potential
Patients with
DRE

150K

CannEpi Treatment Estimate
per patient per annum **£7000³**

Forwards Looking:

- Commercial strategy is a combination of 3rd party distribution supported by in-house Medical Scientific Liaison
- Key wholesaler partners in the UK and Europe: PCCA and Medisonal clinic
- Continuing the work with the 'I am Billy' Foundation and the pathway to NHS RESCAS for Paediatric Refractory Epilepsy
- Dedicated in-house Neurology and Paediatrician collaboration with the European Paediatric Neurology Society⁴
- Initiate CannEpi[®] IND submission to the US FDA



Key points

- CannEpi[®] already sold in the UK, Ireland and Australia
- There are 501 million epilepsy sufferers of which 6.21 million in Europe and the UK

Approximately 30% of generalised seizure epilepsy patients have Refractory Epilepsy aka "Drug-Resistant Epilepsy" (DRE)²

1. <https://www.who.int/news-room/fact-sheets/detail/epilepsy> Fattorusso A, et al. (2021) The Pharmacoresistant Epilepsy

2. Overview on Existant and New Emerging Therapies. Front. Neurol. 12:674483.

3. Alacrita Report 2019

4. <http://dpnsee.org/2019/01/22/treatment-with-medicines-derived-from-cannabis/>

Early Stage Revenue Generating Immunomodulation Treatment for ARDS

CimetrA® delivered 50,000 units to the USA market (Under Special Access) and over 100,000 worldwide, representing over US\$2,500,000 in sales to date.

Results of Preclinical and Clinical program, meeting FDA guidelines, to date:

- Demonstrated **suppression of cytokine storm** in Phase II Clinical Trials¹
- Demonstrated significant reduction in **C-reactive protein (CRP)**, a major inflammatory marker².
- The biological markers in the blood tests further proved a **reduction of inflammation**, enterohepatic involvement, and liver reactant proteins¹.
- Effective **blocking of the IL-32mRNA** expression¹, the pro-inflammatory cytokine related to Autoimmune diseases, lupus, rheumatoid arthritis, inflammatory bowel disease, asthma, and chronic obstructive pulmonary disease¹
- Preclinical studies in rodents (rats, mice) and non-rodent (swine) confirm the safety profile, with no formulation-related toxicity detected³.

1. Data on file – CimetrA in-vitro study

2. Data on file – Interim results – CimetrA Dose Finding Study

3. Data on file – CimetrA pre-clinical study

4. <https://www.mordorintelligence.com/industry-reports/acute-respiratory-distress-syndrome-treatment-market>



Total Addressable
Global Market: £13.6B⁴

Market potential

Cases of severe illness of influenza worldwide¹ **3-5M** Per annum Respiratory deaths **290-650K** Per annum

Potentially, CimetrA® can be used to treat **inflammatory** conditions with **£100B²** total addressable market.

Key points

- Sales strategy is to license and distribute through 3rd party distribution supported by in-house Medical Scientific Liaison
- Key wholesaler partners in the USA and MENA — AMC
- Initiate CimetrA® IND submission to the US FDA

Strategic Milestones

CimetrA® IND submission to the US FDA



1. [https://www.who.int/news-room/fact-sheets/detail/influenza-\(seasonal\)](https://www.who.int/news-room/fact-sheets/detail/influenza-(seasonal))
2. <https://www.precedenceresearch.com/anti-inflammatory-therapeutics-market>

Improving quality of life in Dementia and Alzheimer's

- **55 million people worldwide** suffering from Dementia and Alzheimer's¹
- **CogniCann®** designed as a treatment for the symptoms associated with Dementia and Alzheimer's
- Patients in the Placebo group experienced a deterioration in their condition, compared with the **stable neuropsychiatric profile** of those patients in the treatment group with **CogniCann®**²
- This important finding indicates **improvement in the health** status of the patients and also **the improved quality of life** of the families and caregivers taking care of dementia patients



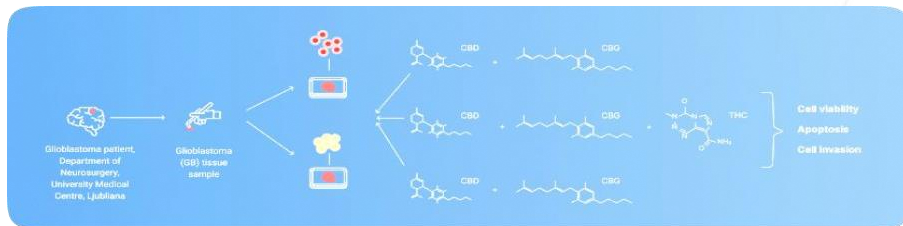
Oromucosal Spray
Pharmaceutical Dosage
Form

1. <https://www.who.int/news-room/fact-sheets/detail/dementia>

2. Data on file : CogniCann Phase II study, Australia

Pre-Clinical Stage Glioblastoma

- Innovative pre-clinical stage drug targeting Glioblastoma Multiforme (GBM) Stem Cells, one of the deadliest forms of brain cancer, with the potential to transform treatment outcomes for this aggressive disease
- An estimated 250,000 new cases of GBM per year worldwide.¹
- The preclinical studies demonstrated that IrniCan® was cytotoxic to Glioblastoma tumor and stem cells, reducing cell viability and inducing, which could establish a new benchmark caspase-dependent cell apoptosis (or cell death in treating GBM by improving safety and efficacy profiles.²



Tablets
Pharmaceutical Dosage
Form

1. <https://www.sciencedaily.com/releases/2024/02/240215113623.htm>

2. Lah, T. . et al. The Cytotoxic Effects of Cannabidiol and Cannabigerol on Glioblastoma Stem Cells May Mostly Involve GPR55 and TRPV1 Signalling. Cancers 2022, 14, 5918T

Pre-Clinical Stage for Chronic Wounds Treatment


- Chronic wounds are a major global health issue, costing billions in treatment annually (e.g., £3.2 billion in the UK ¹ in 2015; \$28.1 to \$96.8 billion in the USA in 2014²). These wounds cause extreme suffering with severe symptoms, often complicated by antibiotic-resistant infections, impenetrable biofilms and deteriorating local tissue health.
- The aim is to develop a nano-formulated product with antimicrobial, biofilm-disrupting, symptom-controlling, and wound-healing properties to address chronic wounds lacking effective treatment.
- Goals of Preclinical Studies:
 - Identify and Select Novel Repurposed Antimicrobial Agents
 - Optimize Combinations for tissue health and relief
 - Enhance Nano-Formulation to maximize therapeutic benefits

Research and Development Product Pipeline

US FDA & EMA Registration Plan

Actively pursuing FDA approval with potential Orphan Drug Designation.



The background features a soft, abstract design with pastel shades of blue, purple, and pink. Large, rounded, overlapping shapes in these colors create a layered effect. A thin, light-colored line curves across the upper left portion of the image. In the upper right corner, there is a small, light blue circle with a subtle gradient and a slight shadow, giving it a three-dimensional appearance.

**Qualified team with versatile
industry experience and expertise**



**Roby
Zomer**

Chief Executive Officer



Roby Zomer is an accomplished executive with extensive experience in the biopharmaceutical and biotech industries. As the CEO of the company, Roby has demonstrated strong leadership in driving strategic growth, managing financial operations, and ensuring compliance with industry-specific standards. He also served as the Chairman of the Board for Graft Polymer, overseeing the implementation of advanced polymer solutions in the Biotech, Automotive and Recycling sectors. Roby's entrepreneurial background includes founding and leading Green City Urban Recycling, a pioneering Israeli company focused on biofuel production. His innovative work in this field contributed to national energy independence initiatives and led to the company's acquisition by Rafael Advanced Defense Systems. With a solid industrial engineering and management foundation, Roby Zomer brings a unique blend of technical and strategic skills to his professional endeavors.



**Yifat
Steuer**

Chief Operating Officer



Yifat is a seasoned executive with over 20 years of experience at GSK, J&J, and various biopharmaceutical and medical device SMEs. With expertise in financial strategy and corporate management, she excels in optimizing cash flows, improving efficiency, and driving large-scale corporate transformations. Yifat holds an MBA in Finance and Strategy and CPA credentials from Deloitte. Yifat leads operational functions, guiding the company's work plans towards sustainable growth. She adeptly manages strategic initiatives, risk mitigation, and corporate relations, significantly boosting company growth and efficiency. Her leadership continues to elevate the company's performance and strengthen its market position.



**Igor
Bluvstein**

Chief Financial Officer



Igor is a seasoned Chief Financial Officer with extensive experience across diverse industries, including digital health, e-commerce, biotechnology, petrochemicals, and medical cannabis. He has held leadership roles as CFO at G Medical Innovations Holdings Ltd., MDD Group of Companies, and an e-commerce retailer. Igor also served as Regional CFO at Frutarom Industries Ltd. and Financial Controller at Mirland Development Corporation PLC, beginning his career as a Senior Auditor at Ernst & Young. He holds a Bachelor of Arts in Accounting and Economics from the Open University in Israel and is a Certified Public Accountant (CPA). Igor leads the finance and compliance department. His international business experience and strong financial management skills make him an exceptional CFO, particularly in publicly traded companies.



**Amir
Polak**

Chief Pharmaceutical
Development Officer



Amir is a seasoned professional with over 18 years of experience in the pharmaceutical and chemical industries. Currently serving as the Pharmaceutical Development Officer at Argent Biopharma LTD, Amir has a strong background in strategic planning, project management, and intellectual property management, with a proven track record of bringing innovative treatments to market. His prior roles include leadership positions at Nano-Dimension Tech, where he spearheaded advanced manufacturing lines for nano-particles, and Green City-Urban Recycling, where he co-founded and led technological advancements. Amir holds an M.Sc. in Organic Chemistry and a B.Sc. in Physical Chemistry from the Hebrew University of Israel. Amir leads the full lifecycle of pharmaceutical product development, from ideation to commercialization.



**Sabina
Suljaković**

Chief Quality and
Commercial Officer



Sabina is an accomplished quality professional with over 10 years of experience, specializing in establishing systems that ensure regulatory compliance and drive commercial success. Sabina led the company to become the only EU GMP-certified entity by JAZMP. As the primary liaison with Health Authorities worldwide, she expertly navigates complex regulatory landscapes, develops strategies for market entry, and leads diverse teams. Her expertise spans regulatory compliance, auditing, production, quality control, and batch certification, with a strong focus on aligning quality initiatives with business goals. Sabina oversees quality, business development, and commercial activities. She is dedicated to fostering continuous improvement and driving both quality and commercial success within the organization.



**Yair
Tal**

Chief Information
Security Officer



With 30 years of extensive experience, Yair Tal specializes in providing comprehensive security solutions. His expertise spans across various aspects of security, ensuring that businesses and organizations operate with confidence in an increasingly complex global environment. He covers security management, data protection, and physical cyber security, from the company's facilities to its online operations and most significantly, patient data. Yair Tal's dedication to security excellence and his broad expertise make him a trusted partner for organizations looking to safeguard their operations and assets.

Medical advisors



Dr. Jonathan Grunfeld

Vice President Medical Development



Dr. Jonathan Grunfeld is a highly experienced neurologist and neuro-oncologist with a career spanning over two decades. He completed his medical degree at Tel Aviv University, followed by a fellowship in neuro-oncology at the prestigious MD Anderson Cancer Center. Dr. Grunfeld has dedicated much of his career to the management of oncological symptoms and palliative care, particularly exploring the therapeutic potential of cannabis in these areas. His extensive experience and innovative approach have equipped him with exceptional clinical observations and unique insights, which he brings to bear in translational initiatives.



Prof. Uri Kramer

Head of Neurology



Professor Uri Kramer is a renowned neurologist, epileptologist, a leading specialist in Israel in the diagnosis and treatment of childhood neurological disorders. Graduated from Medical University in Tel Aviv. Improved his qualifications in childhood epilepsy and clinical neurophysiology in Boston. He is a researcher and creator of several methods in the diagnosis and treatment of epilepsy. He has written over 80 scientific medical papers on pediatric neurology. Created a special EpiLert bracelet with a transmitter that notifies children or adults of an approaching seizure. Member of the European, American and Israeli Associations of Epileptology. He has over 50 years of professional experience.



Dr. Shlomo Sadoun

Strategic Business Development



Dr. Shlomo Sadoun has over 18 years of experience in the pharmaceutical industry. As co-founder and leader of SK-Pharma Group, he has expanded the company into 18 countries, specializing in generic, specialty, and biosimilar products. He also serves as CEO of Arphio, focusing on orphan drugs. Dr. Sadoun has forged strategic alliances with over 200 companies and launched more than 300 products globally, making SK-Pharma one of Israel's fastest-growing pharmaceutical groups. He holds a master's degree in Global Management and a Doctorate in Business Administration with a focus on public health.



Yossi Mograbi

Strategic R/D



Mograbi is a seasoned biotech entrepreneur, having initiated eight projects with top scientists from leading academic and healthcare institutions. Four projects involved NCEs from the Hebrew University of Jerusalem, and another was a drug collaboration with Hadassah Medical Organization. Currently, he's engaged in three projects with Tel-Aviv University, focusing on CSS, COVID-19, and bone applications. As an inventor or co-inventor on seven patents, Yossi has also consulted for Teva, Yissum, and Syqe Medical. He's mentored MBA-Biomed students at the Hebrew University and participated in the SPARK Program. Beyond biotech, Yossi is an expert in dietary supplements, advising companies like Teva and aiding patients with severe rare diseases. His company received two NOFAR grants for cancer and lupus projects, and he assisted with two Kamin grants from Hadasit/Yissum and BGNegev/Hadasit.

Financials¹

PROFIT AND LOSS (USD)	<u>JUN 23- JUL 24*</u>	<u>JUN 22- JUL 23</u>
Revenue	894,263	2,241,621
Gross profit	315,845	959,620
Operating loss	(10,957,390)	(13,149,334)
Loss for the year	(11,523,050)	(13,984,482)

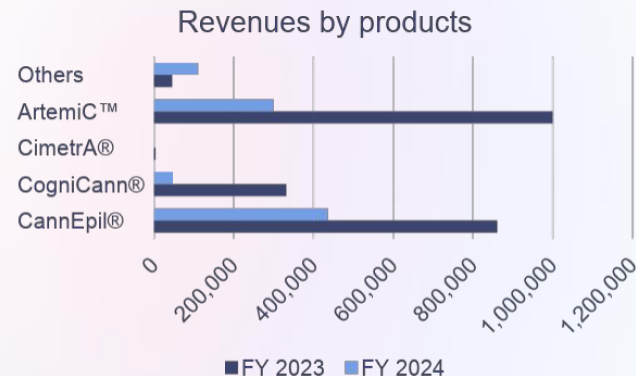
BALANCE SHEET (USD)	<u>JUN 23- JUL 24*</u>	<u>JUN 22-JUL 23</u>
Total assets	7,074,104	6,606,382
Total liabilities	10,778,676	11,921,208
Total equity	(3,704,572)	(5,314,826)

CASH FLOW (USD)	<u>JUN 23- JUL 24</u>	<u>JUN 22- JUL 23</u>
Net cash used in operating activities	(9,511,069)	(7,930,829)
Net cash used in investing activities	(93,448)	401,131
Net cash provided by financing activities	9,913,763	6,440,159
Cash and cash equivalents at the end of the year	468,744	158,694

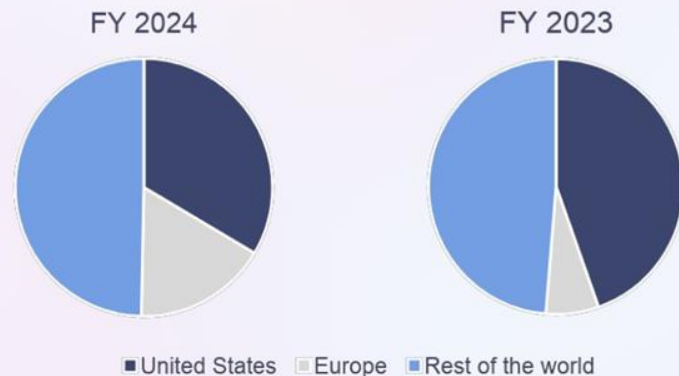
(*) USD 300k revenue reclassification from Sept 2024 due to the timing of invoice

Revenues by products

Sales by products (USD)	JUN 23- JUL 24	JUN 22 – JUL 23
CannEpil®	436,334	860,699
CogniCann®	47,300	331,615
CimetrA®	-	3,862
ArtemiC™	* 300,000	1,000,000
Others	110,629	45,445
SUM	894,263	2,241,621



Sales by GEO region(USD)	JUN 23 – JUL 24	JUN 22 – JUL 23
United States	* 300,000	1,000,000
Europe	149,961	149,627
Rest of the world	444,302	1,091,994



(*) USD 300k revenue reclassification from Sept 2024 due to the timing of invoice



Addresses unmet Medical needs

✉ info@argentbiopharma.com

f [@argentbiopharma](#)

X [@ArgentBioPharma](#)

in [Argent BioPharma Ltd.](#)



[Argent BioPharma](#)